

Ac-(SEQ ID NO: 175) (hereinafter referred to as BCY8157);  
 Ac-(SEQ ID NO: 176) (hereinafter referred to as BCY8158);  
 Ac-(SEQ ID NO: 177) (hereinafter referred to as BCY8161);  
 Ac[B-Ala][Sar10]-(SEQ ID NO: 177) (hereinafter referred to as BCY8278);  
 Ac-(SEQ ID NO: 178) (hereinafter referred to as BCY8162);  
 Ac[B-Ala][Sar10]-(SEQ ID NO: 178) (hereinafter referred to as BCY8277);  
 Ac-(SEQ ID NO: 179) (hereinafter referred to as BCY8163);  
 Ac-[B-Ala][Sar10]-(SEQ ID NO: 179) (hereinafter referred to as BCY8276);  
 [B-Ala][Sar10]-(SEQ ID NO: 179) (hereinafter referred to as BCY8269);  
 Ac-(SEQ ID NO: 180) (hereinafter referred to as BCY8174);  
 Ac-(SEQ ID NO: 181) (hereinafter referred to as BCY8175);  
 Ac-(SEQ ID NO: 182) (hereinafter referred to as BCY8176);  
 Ac-(SEQ ID NO: 183) (hereinafter referred to as BCY8177);  
 Ac-(SEQ ID NO: 184) (hereinafter referred to as BCY8178);  
 Ac-(SEQ ID NO: 185) (hereinafter referred to as BCY8180);  
 Ac-(SEQ ID NO: 186) (hereinafter referred to as BCY8181);  
 Ac-(SEQ ID NO: 187) (hereinafter referred to as BCY8182);  
 Ac-(SEQ ID NO: 188) (hereinafter referred to as BCY8183);  
 Ac-(SEQ ID NO: 189) (hereinafter referred to as BCY8184);  
 [B-Ala][Sar10]-(SEQ ID NO: 189) (hereinafter referred to as BCY8235);  
 Ac-(SEQ ID NO: 190) (hereinafter referred to as BCY8185);  
 Ac-(SEQ ID NO: 191) (hereinafter referred to as BCY8186);  
 Ac-(SEQ ID NO: 192) (hereinafter referred to as BCY8187);  
 Ac-(SEQ ID NO: 193) (hereinafter referred to as BCY8188);  
 Ac-(SEQ ID NO: 194) (hereinafter referred to as BCY8189);  
 Ac-(SEQ ID NO: 195) (hereinafter referred to as BCY8191);  
 Ac-(SEQ ID NO: 196) (hereinafter referred to as BCY8192);  
 Ac-(SEQ ID NO: 197) (hereinafter referred to as BCY8193);  
 Ac-(SEQ ID NO: 198) (hereinafter referred to as BCY8194);  
 Ac-(SEQ ID NO: 199) (hereinafter referred to as BCY8211);  
 Ac-(SEQ ID NO: 200) (hereinafter referred to as BCY8212);  
 Ac-(SEQ ID NO: 201) (hereinafter referred to as BCY8213);

Ac-(SEQ ID NO: 202) (hereinafter referred to as BCY8214);  
 [B-Ala][Sar10]-(SEQ ID NO: 202) (hereinafter referred to as BCY8231);  
 Ac-(SEQ ID NO: 203) (hereinafter referred to as BCY8215);  
 Ac[B-Ala][Sar10]-(SEQ ID NO: 208) (hereinafter referred to as BCY8279);  
 Ac[B-Ala][Sar10]-(SEQ ID NO: 209) (hereinafter referred to as BCY8280);  
 [B-Ala][Sar10]-(SEQ ID NO: 209) (hereinafter referred to as BCY8273);  
 Ac[B-Ala][Sar10]-(SEQ ID NO: 210) (hereinafter referred to as BCY8281);  
 Ac-(SEQ ID NO: 211) (hereinafter referred to as BCY8831);  
 [B-Ala][Sar10]-(SEQ ID NO: 212) (hereinafter referred to as BCY8238);  
 (SEQ ID NO: 215) (hereinafter referred to as BCY11415);  
 [PYA][B-Ala][Sar10]-(SEQ ID NO: 215) (hereinafter referred to as BCY11942); and  
 (SEQ ID NO: 216) (hereinafter referred to as BCY11414).

**11.** The peptide ligand as defined in any one of claims **1** to **10**, wherein the molecular scaffold is selected from 1,1',1''-(1,3,5-triazinane-1,3,5-triyl)triprop-2-en-1-one (TATA).

**12.** The peptide ligand as defined in any one of claims **1** to **11**, wherein the pharmaceutically acceptable salt is selected from the free acid or the sodium, potassium, calcium, ammonium salt.

**13.** The peptide ligand as defined in any one of claims **1** to **12**, wherein the Nectin-4 is human Nectin-4.

**14.** A drug conjugate comprising a peptide ligand as defined in any one of claims **1** to **13**, conjugated to one or more effector and/or functional groups.

**15.** The drug conjugate as defined in claim **14**, conjugated to one or more cytotoxic agents.

**16.** The drug conjugate as defined in claim **15**, wherein said cytotoxic agent is selected from MMAE or DM1, in particular MMAE.

**17.** The drug conjugate as defined in claim **16**, wherein the cytotoxic agent is MMAE and said conjugate additionally comprises a linker selected from: -PABC-Cit-Val-Glutaryl- or -PABC-cyclobutyl-Ala-Cit-pAla-, such as -PABC-Cit-Val-Glutaryl-, wherein PABC represents p-aminobenzylcarbamate.

**18.** The drug conjugate as defined in claim **16**, wherein the cytotoxic agent is DM1 and said conjugate additionally comprises a linker which is -SPDB-(SO<sub>3</sub>H)—, wherein SPDB represents N-succinimidyl 3-(2-pyridyldithio)propionate.

**19.** The drug conjugate as defined in any one of claims **14** to **18**, which is selected from any one of: BCY7683, BCY7825, BCY7826, BCY8245, BCY8253, BCY8254, BCY8255, BCY8549, BCY8550, BCY8783 and BCY8784, such as: BCY7683, BCY7825, BCY7826, BCY8245, BCY8253, BCY8254, BCY8255, BCY8783 and BCY8784, in particular BCY8245.

**20.** A pharmaceutical composition which comprises the peptide ligand of any one of claims **1** to **13** or the drug conjugate of any one of claims **14** to **19**, in combination with one or more pharmaceutically acceptable excipients.